

GenCore version 5.1.5
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OM nucleic - nucleic search, using sw model

Run on: June 1, 2003, 14:56:34 ; Search time 466.436 Seconds
(without alignments)
10776.324 Million cell updates/sec

Title: US-09-625-573-1
Perfect score: 2232
Sequence: 1 GGATTGAACAGGACGCAATT.....TATAACTATGTCATATAAAG 2232

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N_Geneseq_101002.*
1: /SID22/cgdata/geneseq/geneseq-emb1/NA1980.DAT.*
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3: /SID22/cgdata/geneseq/geneseq-emb1/NA1982.DAT.*
4: /SID22/cgdata/geneseq/geneseq-emb1/NA1983.DAT.*
5: /SID22/cgdata/geneseq/geneseq-emb1/NA1984.DAT.*
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22: /SID22/cgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
23: /SID22/cgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
24: /SID22/cgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	2232	100.0	2232	16 AAQ96297	Human monocyte che
2	1250.8	56.0	143068	21 AAF21105	Human low adenosin
3	1250.8	56.0	143068	21 AAF21272	Human low adenosin
4	1250.8	56.0	143068	21 AAA34983	Human adenosine re
5	1250.8	56.0	143068	21 AAA35150	Human adenosine re
6	1250.8	56.0	143068	21 ABL68124	Ovary cancer relat
7	1250.8	56.0	149412	21 AAA35151	Human adenosine re
8	1250.8	56.0	152740	21 AAF21273	Human low adenosin
9	980	43.9	1979	16 AAQ96298	Human monocyte che

10	941	42.2	1083	22	AA512140	Human wild-type CC
11	939.4	42.1	1083	22	AA512139	Human CCR2-641 pol
12	937.8	42.0	1083	18	AA796976	Human monocyte che
13	937.8	42.0	1083	23	AB197976	Non-endogenous hum
14	668.2	29.9	10528	24	ABL32335	Human immune syste
15	639.8	28.7	10528	24	ABL32334	Human immune syste
16	635.6	28.5	1059	18	AA785163	Macaque chemokine
17	635.6	28.5	1059	22	AA165463	Nucleotide sequenc
18	632.6	28.3	1056	22	AA165463	Human G-protein ch
19	632.6	28.3	1056	22	AA165463	Human G-protein ch
20	632.6	28.3	1056	22	AA165463	DNA encoding human
21	632.6	28.3	1059	19	AAV23992	Human CC-CKR5 codi
22	632.6	28.3	1059	24	ABA97319	Human chemokine (C
23	632.6	28.3	1071	20	AAV84125	HIV-1 co-receptor
24	632.6	28.3	1225	19	AA76920	DNA encoding human
25	632.6	28.3	1225	24	ABA02317	Human CC chemokine
26	632.6	28.3	1376	22	AAH26903	Human HIV-1 co-rec
27	632.6	28.3	1414	22	AAF26390	Human HDGNR10 CDNA
28	632.6	28.3	1477	18	AA790117	CDNA for human CCR
29	632.6	28.3	1477	22	AAF87099	Human CCR5 CDNA se
30	632.6	28.3	3383	18	AA785161	Human chemokine re
31	632.6	28.3	3383	21	AA721271	Human low adenosin
32	632.6	28.3	3383	21	AAA35149	Human adenosine re
33	632.6	28.3	3383	22	AAH08577	Human chemokine re
34	632.6	28.3	5674	20	AAZ24738	Human chemokine re
35	632.6	28.3	9141	24	ABA97318	Human chemokine (C
36	631.2	28.3	1376	20	AAV84126	Human co-receptor
37	631	28.3	1225	24	ABA02318	HIV-1 co-receptor
38	629.4	28.2	1059	23	AB197978	Non-endogenous hum
39	629.4	28.2	1414	18	AA794042	Human G-protein ch
40	629.4	28.2	1414	21	AAZ91481	Human G-protein ch
41	629.4	28.2	1414	22	AAZ91481	Human G-protein ch
42	629.4	28.2	1414	22	AAZ91481	Human G-protein ch
43	629.4	28.2	1414	24	ABK51853	DNA encoding human
44	621.4	27.8	1255	19	AA76919	DNA encoding human
45	612.6	27.4	1344	20	AAV84159	HIV-1 co-receptor

ALIGNMENTS

RESULT 1
AAQ96297
ID AAQ96297 standard; cDNA; 2232 BP.

AC AAQ96297;

XX 29-DEC-1995 (first entry)

DE Human monocyte chemoattractant protein-1 receptor MCP-1RA.

XX Monocyte chemoattractant protein-1 receptor; MCR-1R; chemokine; ss.

OS Homo sapiens.

XX Key Location/Qualifiers
FH CDS 40..1161
FT /*tag= a

XX WO9519436-A.

XX 20-JUL-1995.

XX 11-JAN-1995; 95WO-US00476.

XX 13-JAN-1994; 94US-0182962.

XX (REGC) UNIV CALIFORNIA.

XX Charo I, Coughlin S;

XX WPI; 1995-263866/34.

XX P-PSDB; AAR9165.

DR

XX

PT DNA encoding monocyte chemo-attractant protein-1 receptor - used partic.
PT for identifying antagonists and for treating diseases characterised by
XX monocytic infiltrates

PS Disclosure; Fig 1; 84pp; English.

XX To identify and clone new members of the chemokine receptor gene
CC family, degenerate oligo primers were designed correspond. to the
CC conserved sequences R79167 in the second and R79168 in the third
CC transmembrane domains of the MIP-lalpha/RANTES receptor, the IL-8
CC receptors and the HUMSTRS orphan receptor (GenBank Accession #99293.
CC The degenerate oligo incorporating EcoRI and XhoI sites at their 5',
CC ends are Q96299 and Q96300. Amplification of cDNA derived from MM6
CC cells with the primers yielded a number of PCR products. One cDNA
CC appeared to encode a novel protein. To obtain a full-length version
CC of this clone, a MM6 cDNA library was constructed in pPROG and probed
CC with the PCR product. A 2.1 kb cDNA clone was obtained. Analysis of
CC additional clones in the MM6 cDNA library revealed a second
CC sequence that was identical to the 2.1 kb cDNA sequence first obtained.
CC from the 5' UTR through the putative seventh transmembrane domain
CC but contained a different cytoplasmic tail. The second sequence
CC appears to represent alternative splicing of the carboxyl-terminal
CC tail of the MCP-1R protein. The two sequences are denoted MCP-1RA
CC and MCP-1RB (see Q96297/R79165 & Q96298/R79166). Active mature
CC MCP-1RA has a predicted mol. wt. of about 42,000 daltons. MCP-1RB
CC has a mol. wt. of about 41,000 daltons.

XX Sequence 2232 BP; 602 A; 464 C; 508 G; 658 T; 0 other;

Query Match 100.0%; Score 2232; DB 16; Length 2232;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 2232; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGATTGAACAGGAGCGATTTCCCGAGTACATCCACAAACATGCTGCCACATCTCGTTCT 60
DB 1 GGATTGAACAGGAGCGATTTCCCGAGTACATCCACAAACATGCTGCCACATCTCGTTCT 60
QY 61 CGGTTTATCAGAAATACCAACGAGAGCGGTGAAGAGTCCACACCTTTTGGATTATGAT 120
DB 61 CGGTTTATCAGAAATACCAACGAGAGCGGTGAAGAGTCCACACCTTTTGGATTATGAT 120
QY 121 TACGGTCTCCCTGTCATAAATTTGACGTGAAGCAAAATTTGGGGCCCACTCTCGCTCG 180
DB 121 TACGGTCTCCCTGTCATAAATTTGACGTGAAGCAAAATTTGGGGCCCACTCTCGCTCG 180
QY 181 CTCCTACTCGCTGGTTCATCTTTGGTGTGGGCAACATGCTGGTCTCTCATCTTA 240
DB 181 CTCCTACTCGCTGGTTCATCTTTGGTGTGGGCAACATGCTGGTCTCTCATCTTA 240
QY 241 ATAACTGCAAAAGCTGAAGTGTGACTGACATTTACCTGCTCAACCTGGCCATCTCT 300
DB 241 ATAACTGCAAAAGCTGAAGTGTGACTGACATTTACCTGCTCAACCTGGCCATCTCT 300
QY 301 GATCTGCTTTTCTATTACTCTCCCATTTGGGCTCACTCTGCTGCAAAATGAGTGGGTC 360
DB 301 GATCTGCTTTTCTATTACTCTCCCATTTGGGCTCACTCTGCTGCAAAATGAGTGGGTC 360
QY 361 TTTGGGAATGCAATGTCAAAATTTACAGGGCTGTATCATCATCGGTTATTTGGCCGA 420
DB 361 TTTGGGAATGCAATGTCAAAATTTACAGGGCTGTATCATCATCGGTTATTTGGCCGA 420
QY 421 ATCTTCTTCATATCCCTCTGACAAATCGATAGATACCTGGCTATTGCTCATGCTGTGTT 480
DB 421 ATCTTCTTCATATCCCTCTGACAAATCGATAGATACCTGGCTATTGCTCATGCTGTGTT 480
QY 481 GCTTTAAAGCCAGGAGGTCACCTTTGGGGTGTGATCAAGTGTGATCACTGGTTGGTG 540
DB 481 GCTTTAAAGCCAGGAGGTCACCTTTGGGGTGTGATCAAGTGTGATCACTGGTTGGTG 540
QY 541 GCTGTCTTTGCTTCTGTCCAGGAATCATCTTTACTAAATGCCAGAGAGATCTGTT 600
DB 541 GCTGTCTTTGCTTCTGTCCAGGAATCATCTTTACTAAATGCCAGAGAGATCTGTT 600

QY 601 TATGCTGTGGCCCTTATTTTCCAGGAGGATGAATAATTTCCACACAATAATGAGGAAC 660
DB 601 TATGCTGTGGCCCTTATTTTCCAGGAGGATGAATAATTTCCACACAATAATGAGGAAC 660
QY 661 ATTTTGGGGCTGCTCCTCGCTGCCTCATCATGCTCATCTGCTACTCGGGAATCCTGAAA 720
DB 661 ATTTTGGGGCTGCTCCTCGCTGCCTCATCATGCTCATCTGCTACTCGGGAATCCTGAAA 720
QY 721 ACCCTGCTTCCGCTGTGCGAAGAGAGGATAGGCGATAGGCGAGTGCATCTTCCACC 780
DB 721 ACCCTGCTTCCGCTGTGCGAAGAGAGGATAGGCGATAGGCGAGTGCATCTTCCACC 780
QY 781 ATCATGATGTTTACTTCTTCTGAGTCCCTATAACATTTGCTATCTTCTCTGAAACACC 840
DB 781 ATCATGATGTTTACTTCTTCTGAGTCCCTATAACATTTGCTATCTTCTCTGAAACACC 840
QY 841 TTCCAGGAATTTCTCGGCTGAGTAACTGTGAAGCACCAGTCACTGGACCAAGCCACG 900
DB 841 TTCCAGGAATTTCTCGGCTGAGTAACTGTGAAGCACCAGTCACTGGACCAAGCCACG 900
QY 901 CAGGTGACAGAGACTTCTGGGATGACTCACTGCTGCATCAATCCCATCTATCTATGCTTC 960
DB 901 CAGGTGACAGAGACTTCTGGGATGACTCACTGCTGCATCAATCCCATCTATCTATGCTTC 960
QY 961 GTTGGGGAAGTTTCAGAAAGCTTTTTCACATAGCTCTTGGCTGTAGGATTGCCCCACTC 1020
DB 961 GTTGGGGAAGTTTCAGAAAGCTTTTTCACATAGCTCTTGGCTGTAGGATTGCCCCACTC 1020
QY 1021 CAAAACACAGTGTGTGGAGTCCAGGAGTGAGACAGGAGAAAGAAATGTGAAAGTGACTACA 1080
DB 1021 CAAAACACAGTGTGTGGAGTCCAGGAGTGAGACAGGAGAAAGAAATGTGAAAGTGACTACA 1080
QY 1081 CAAGGACTCTCTGATGCTGTGGAAAGGAAGTCAATTTGGCAGAGCCCCCTGAAGCCAGT 1140
DB 1081 CAAGGACTCTCTGATGCTGTGGAAAGGAAGTCAATTTGGCAGAGCCCCCTGAAGCCAGT 1140
QY 1141 CTTGAGCAAAAGAGGAGCCTTAGAGACAGAAATGACAGATCTCTGCTTTGGAAATCACA 1200
DB 1141 CTTGAGCAAAAGAGGAGCCTTAGAGACAGAAATGACAGATCTCTGCTTTGGAAATCACA 1200
QY 1201 CGTCTGCTTTCACAGATGTGTGATTTACAGTGTGAATTTGGTGTCTACGTACCCAGCA 1260
DB 1201 CGTCTGCTTTCACAGATGTGTGATTTACAGTGTGAATTTGGTGTCTACGTACCCAGCA 1260
QY 1261 GGAAGGCTGAGAGGAGAGACTCCAGTGGTGGTGGAAACAGATTTTCCAAACTACCT 1320
DB 1261 GGAAGGCTGAGAGGAGAGACTCCAGTGGTGGTGGAAACAGATTTTCCAAACTACCT 1320
QY 1321 TCAGTTCCTCATTTTGAATACAGGATAGAGTTCAGACTTTTTTAAATAGTAAAT 1380
DB 1321 TCAGTTCCTCATTTTGAATACAGGATAGAGTTCAGACTTTTTTAAATAGTAAAT 1380
QY 1381 AAAATTAAGCTCAAAACTGCAACTTGAATGTGTTAAAGAGTGTAGTTGAGTTGCTPAT 1440
DB 1381 AAAATTAAGCTCAAAACTGCAACTTGAATGTGTTAAAGAGTGTAGTTGAGTTGCTPAT 1440
QY 1441 CATGTCAAAAGCTGAAAATGCTGTTAGTACAGAGATAATTTAGCTTTTGGAGCTTAA 1500
DB 1441 CATGTCAAAAGCTGAAAATGCTGTTAGTACAGAGATAATTTAGCTTTTGGAGCTTAA 1500
QY 1501 ATTTTTCAGAGGTTGTTGTTGGGAGACTGCTGAGTCAACCCCAATAGTTGTTGATGGC 1560
DB 1501 ATTTTTCAGAGGTTGTTGTTGGGAGACTGCTGAGTCAACCCCAATAGTTGTTGATGGC 1560
QY 1561 AGGAGTTGGAAGTGTGTGATCTGTGGGACATTTAGCTATGCTGCATCGACATCTAAGTA 1620
DB 1561 AGGAGTTGGAAGTGTGTGATCTGTGGGACATTTAGCTATGCTGCATCGACATCTAAGTA 1620
QY 1621 ATGATGCTGTTTGAATACAGATATACGCTCCATCGCTGTGATCTCAGCTGGATCTCCATT 1680
DB 1621 ATGATGCTGTTTGAATACAGATATACGCTCCATCGCTGTGATCTCAGCTGGATCTCCATT 1680

QY 1681 CTCTCAGGCTGCTGCCAAAGCCCTTTGTTGTTTGTATCATTTATGAAGTCATGC 1740
Db 1681 CTCTCAGGCTGCTGCCAAAGCCCTTTGTTGTTTGTATCATTTATGAAGTCATGC 1740
QY 1741 GTTTAATCACATTCAGTGTTCAGTGTTCGACATGTCCTTGATGTCATATTTGTTCC 1800
Db 1741 GTTTAATCACATTCAGTGTTCAGTGTTCGACATGTCCTTGATGTCATATTTGTTCC 1800
QY 1801 CTAATTTGCCAGTGGGAACCTCTAAATCAAAATTTGCTTCTAATCAAAAGCTTTTAAACCC 1860
Db 1801 CTAATTTGCCAGTGGGAACCTCTAAATCAAAATTTGCTTCTAATCAAAAGCTTTTAAACCC 1860
QY 1861 ATTGTAAGATGAAGTGGAGAGCTCCCTGAAGTAAGCAAGACTTTCTCTTAGT 1920
Db 1861 ATTGTAAGATGAAGTGGAGAGCTCCCTGAAGTAAGCAAGACTTTCTCTTAGT 1920
QY 1921 CGAGCCAAAGTTAAGATGTTCTTATGTTGCCAGTGTGTTCTGATGTCGATCAAGCAAG 1980
Db 1921 CGAGCCAAAGTTAAGATGTTCTTATGTTGCCAGTGTGTTCTGATGTCGATCAAGCAAG 1980
QY 1981 AAACACTGGGCTTCTAGAACAGGCAACTTTGGGAAGTGGGAGTGGGAGTGGGAGTGGG 2040
Db 1981 AAACACTGGGCTTCTAGAACAGGCAACTTTGGGAAGTGGGAGTGGGAGTGGGAGTGGG 2040
QY 2041 TCTACTTTCCAGCCCATGCTTAAAGAGGTTTCAGAAAGAGTGGGAGTGGGAGTGGGAGTGGG 2100
Db 2041 TCTACTTTCCAGCCCATGCTTAAAGAGGTTTCAGAAAGAGTGGGAGTGGGAGTGGGAGTGGG 2100
QY 2101 TTTACCTTCATATATTTGATGATCCTTAATGAATGCATATAAATGTTAAGTTGATGTTGA 2160
Db 2101 TTTACCTTCATATATTTGATGATCCTTAATGAATGCATATAAATGTTAAGTTGATGTTGA 2160
QY 2161 TGAATGTAATTAATCTTTTAACTATGATTTGGAATAAATGTTGGAATAAATGTTGGAATAA 2220
Db 2161 TGAATGTAATTAATCTTTTAACTATGATTTGGAATAAATGTTGGAATAAATGTTGGAATAA 2220
QY 2221 TGTGTATAAAG 2232
Db 2221 TGTGTATAAAG 2232

RESULT 2
AAF21105
ID AAF21105 standard; DNA; 143068 BP.
XX AAF21105;
AC AAF21105;
XX AAF21105;
DT 14-MAR-2001 (first entry)
XX Human low adenine antisense oligonucleotide related sequence #2672.

Low adenine antisense oligonucleotide; phosphorothioate; allergy;
human; airway disorder; bronchoconstriction; lung inflammation;
surfactant depletion; respiratory; bronchodilator; antiinflammatory;
immunosuppressive; antiasthmatic; analgesic; hypotensive; cytostatic;
respiratory obstruction; pulmonary obstruction; impeded respiration;
surfactant hypoproduction; pulmonary vasoconstriction; asthma; RDS;
respiratory distress syndrome; pain; cystic fibrosis; allergic rhinitis;
pulmonary hypertension; emphysema; pulmonary transplantation rejection;
chronic obstructive pulmonary disease; pulmonary infection; bronchitis;
cancer; ss.

XX Homo sapiens.
XX WO200062736-A2.
XX 26-OCT-2000.
XX 24-MAR-2000; 2000WO-US08020.
XX 06-APR-1999; 99US-0127958.
XX (UYEC-) UNIV EAST CAROLINA.

PA (NYCE/) NYCE J W.
XX NYCE JW;
PI WPI; 2000-679539/66.
XX Low adenine (A) content antisense oligonucleotides which do not
PT trigger adenine receptors during metabolism, useful e.g. for treating
PT cancers and respiratory obstructions -
XX Disclosure: Page 924-957; 1592pp; English.
XX The present invention describes low adenine (A) content antisense
CC oligonucleotides and compositions (I) comprising them. In the antisense
CC oligonucleotides the A is replaced by a 'Universal' or alternative base.
CC (I) can have respiratory, bronchodilator, antiinflammatory, analgesic,
CC immunosuppressive, antiasthmatic, hypotensive and cytostatic activities.
CC The antisense oligonucleotides and (I) can be used to down-regulate the
CC expression and or activity of target polypeptides associated with
CC lung/respiratory disorders and malignancies, such as stimulating and
CC activating peptide factors and transmitters, transcription factors and
CC immunoglobulins and antibodies, antibody receptors, cytokines and
CC chemokines, endogenously produced specific and non-specific enzymes,
CC binding proteins, adenosine receptors, bradykinin receptors, central
CC chemokine receptors, adenosine receptors, bradykinin receptors, central
CC nervous system (CNS) and peripheral nervous and non-nervous system
CC receptors, CNS and peripheral nervous and non-nervous system peptide
CC transmitters, defensins, growth factors, vasoactive peptides and
CC receptors, binding proteins and malignancy associated proteins. The
CC antisense oligonucleotides may be used in this way to treat disorders
CC including respiratory obstruction (especially pulmonary obstruction
CC and/or bronchoconstriction) and/or lung inflammation, allergy(ies)
CC and/or surfactant hypoproduction which are associated with a disease or
CC condition selected from pulmonary vasoconstriction, inflammation,
CC allergies, asthma, impeded respiration, respiratory distress syndrome
CC (RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary
CC hypertension, emphysema, chronic obstructive pulmonary disease (COPD),
CC pulmonary transplantation rejection, pulmonary infections, bronchitis,
CC and/or cancer. AAF18434 to AAF21543 represent human polynucleotide
CC fragments and antisense oligonucleotides used in the exemplification of
CC the present invention.
XX SQ Sequence 143068 BP; 41194 A; 30122 C; 32403 G; 39349 T; 0 other;

Query Match 56.0%; Score 1250.8; DB 21; Length 143068;
Best Local Similarity 99.8%; Pred. No. 0;
Matches 1252; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 979 AGCCTTTTTCACATAGCTCTTGGCTGTAGGATGCCCCACTCCAAAACCCAGTGTGGA 1038
Db 48253 AGCCTTTTTCACATAGCTCTTGGCTGTAGGATGCCCCACTCCAAAACCCAGTGTGGA 48312
QY 1039 GTTCCAGGAGTACAGCAGGAGAAAGTGTGAAGTGTACAGGACTCCTCGATGGT 1098
Db 48313 GTTCCAGGAGTACAGCAGGAGAAAGTGTGAAGTGTACAGGACTCCTCGATGGT 48372
QY 1099 CGTGGAAAAGAAAGTCAATTTGGCAGAGCCCTTCAAGCCAGTCTTCAGGACAAAAGGA 1158
Db 48373 CGTGGAAAAGAAAGTCAATTTGGCAGAGCCCTTCAAGCCAGTCTTCAGGACAAAAGGA 48432
QY 1159 GCCTAGAGACAGAAATGACAGATCTCTGCTTTGGAAATCAACAGTGTGGCTTCAGATG 1218
Db 48433 GCCTAGAGACAGAAATGACAGATCTCTGCTTTGGAAATCAACAGTGTGGCTTCAGATG 48492
QY 1219 TGTGATTCACAGTGTGAATCTTGGTGTCTAGTTCACAGGAGGAGGAGGAGAG 1278
Db 48493 TGTGATTCACAGTGTGAATCTTGGTGTCTAGTTCACAGGAGGAGGAGGAGAG 48552
QY 1279 AGACTCCAGCTGGGTGGAAAACAGTATTTTCCAAACTTACCTTCAGTTCCTCATTTTG 1338
Db 48553 AGACTCCAGCTGGGTGGAAAACAGTATTTTCCAAACTTACCTTCAGTTCCTCATTTTG 48612
QY 1339 AATACAGGCATAGAGTTTCAGACTTTTAAATAGTAAAAATAAATTAAGCTGAAAC 1398

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Db 48613 AATACAGGCTAGAGTTACAGCTTTTAAATAGTAAAAATAAATTAAGCTGAAAC 48672
Qy 1399 TCGAACTTTGTAATGTGTAAGAGTTAGTTGAGTTGCTATCATGTCAACAGTGAAAT 1458
Db 48673 TCGAACTTTGTAATGTGTAAGAGTTAGTTGAGTTGCTATCATGTCAACAGTGAAAT 48732
Qy 1459 GCTGTATTAGTACAGAGATAATCTAGCTTTGAGCTTAAAGAAATTTGAGCAGGTGAT 1518
Db 48733 GCTGTATTAGTACAGAGATAATCTAGCTTTGAGCTTAAAGAAATTTGAGCAGGTGAT 48792
Qy 1519 GTTTGGGAGACTGCTGAGTCAACCAATAGTTGCTGATTTGAGCAGGAGTTGGAAGTGTG 1578
Db 48793 GTTTGGGAGACTGCTGAGTCAACCAATAGTTGCTGATTTGAGCAGGAGTTGGAAGTGTG 48852
Qy 1579 ATCTGTGGGACATATAGCTATGTCATGCAGCATCTAAGTAATGATGCTTTGAATCA 1638
Db 48853 ATCTGTGGGACATATAGCTATGTCATGCAGCATCTAAGTAATGATGCTTTGAATCA 48912
Qy 1639 CAGTATAGCTCCATCGCTGTCATCTCAGCTGGATCTCCATCTCFCAGGCTTGTGCCA 1698
Db 48913 CAGTATAGCTCCATCGCTGTCATCTCAGCTGGATCTCCATCTCFCAGGCTTGTGCCA 48972
Qy 1699 AAAGCTTTTGTGTTTGTGTTGATCATTAATGAAGTATGCTTTAATCAATTCAGT 1758
Db 48973 AAAGCTTTTGTGTTTGTGTTGATCATTAATGAAGTATGCTTTAATCAATTCAGT 49032
Qy 1759 GTTTCAGTCTCGCAGATGCTTGTGCTCATATGTTGCTTCCCTAATTTGCCAGTGGAA 1818
Db 49033 GTTTCAGTCTCGCAGATGCTTGTGCTCATATGTTGCTTCCCTAATTTGCCAGTGGAA 49092
Qy 1819 CTCCTAATCAATTTGGCTTCTAATCAAGCTTTTAAACCTATTTGGTAAAGATGGAG 1878
Db 49093 CTCCTAATCAATTTGGCTTCTAATCAAGCTTTTAAACCTATTTGGTAAAGATGGAG 49152
Qy 1879 GTGAGAGCTCCCTGAGTAAAGCAAGACTTTCCTTATGTCGAGCAGCAAGTAAAGATG 1938
Db 49153 GTGAGAGCTCCCTGAGTAAAGCAAGACTTTCCTTATGTCGAGCAGCAAGTAAAGATG 49212
Qy 1939 TTTCTATGTTCCAGTGTGTTCTGATCTGATGCAAGCAAGAACTGGGCTTCTAGA 1998
Db 49213 TTTCTATGTTCCAGTGTGTTCTGATCTGATGCAAGCAAGAACTGGGCTTCTAGA 49272
Qy 1999 ACCAGCAACTTGGGAAGTCCAGTCCAGCTGAGTATGCTGCTACCTTACCTTCAATATTT 2058
Db 49273 ACCAGCAACTTGGGAAGTCCAGTCCAGCTGAGTATGCTGCTACCTTACCTTCAATATTT 49332
Qy 2059 GGCTAAAGAGGTTTTCAGAAAGAGTGGGACAGCAGAGAACTTTCACCTTCAATATTT 2118
Db 49333 GGCTAAAGAGGTTTTCAGAAAGAGTGGGACAGCAGAGAACTTTCACCTTCAATATTT 49392
Qy 2119 GTATGATCCTTAATCAATGATCAATGATTAAGTTGATGGTGAATGTAATACTGTT 2178
Db 49393 GTATGATCCTTAATCAATGATCAATGATTAAGTTGATGGTGAATGTAATACTGTT 49452
Qy 2179 TTTTAACTATGATTTGGAATAAATAATCAATGCTATTAACATGTTGATAAAG 2232
Db 49453 TTTTAACTATGATTTGGAATAAATAATCAATGCTATTAACATGTTGATAAAG 49506
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RESULT 3

AAAF21272
ID AAF21272 standard; DNA; 143068 bp.

XX AAF21272;

XX AAF21272;

DT 14-MAR-2001 (first entry)

XX Human low adenosine antisense oligonucleotide related sequence #2839.

DE Low adenosine antisense oligonucleotide; phosphorothioate; allergy;
XX human; airway disorder; bronchoconstriction; lung inflammation;
KW surfactant depletion; respiratory; bronchodilator; antiinflammatory;
KW

immunosuppressive; antiasthmatic; analgesic; hypotensive; cytostatic;
respiratory obstruction; pulmonary obstruction; impeded respiration;
surfactant hypoproduction; pulmonary vasoconstriction; asthma; RDS;
respiratory distress syndrome; pain; cystic fibrosis; allergic rhinitis;
pulmonary hypertension; emphysema; pulmonary transplantation rejection;
chronic obstructive pulmonary disease; pulmonary infection; bronchitis;
cancer; ss.
XX
OS Homo sapiens.
PN W0200062736-A2.
XX 26-OCT-2000.
XX
PE 24-MAR-2000; 2000WO-US08020.
XX
PR 06-APR-1999; 99US-0127958.
XX
PA (UYEC-) UNIV EAST CAROLINA.
XX (NYCE/) NYCE J W.
XX Nyce JW;
XX WPI; 2000-679539/66.

Low adenosine (A) content antisense oligonucleotides which do not
trigger adenosine receptors during metabolism, useful e.g. for treating
cancers and respiratory obstructions -
Disclosure; Page 1186-1219; 1592pp; English.

The present invention describes low adenosine (A) content antisense
oligonucleotides and compositions (I) comprising them. In the antisense
oligonucleotides the A is replaced by a 'universal' or alternative base.
(I) can have respiratory, bronchodilator, antiinflammatory, analgesic,
immunosuppressive, antiasthmatic, hypotensive and cytostatic activities.
The antisense oligonucleotides and (I) can be used to down-regulate the
expression and/or activity of target polypeptides associated with
lung/respiratory disorders and malignancies, such as stimulating and
activating peptide factors and transmitters, transcription factors,
immunoglobulins and antibodies, antibody receptors, cytokines and
chemokines, endogenously produced specific and non-specific enzymes,
binding proteins, adhesion molecules and their receptors, cytokine and
chemokine receptors, adenosine receptors, bradykinin receptors, central
nervous system (CNS) and peripheral nervous and non-nervous system
receptors, CNS and peripheral nervous and non-nervous system peptide
transmitters, defensins growth factors, vasoactive peptides and
receptors, binding proteins and malignancy associated proteins. The
antisense oligonucleotides may be used in this way to treat disorders
including respiratory obstruction (especially pulmonary obstruction
and/or bronchoconstriction) and/or lung inflammation, allergy(ies)
and/or surfactant hypoproduction which are associated with a disease or
condition selected from pulmonary vasoconstriction, inflammation,
allergies, asthma, impeded respiration, respiratory distress syndrome
(RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary
hypertension, emphysema, chronic obstructive pulmonary disease (COPD),
pulmonary transplantation rejection, pulmonary infections, bronchitis,
and/or cancer. AAF18434 to AAF21543 represent human polynucleotide
fragments and antisense oligonucleotides used in the exemplification of
the present invention.

Sequence 143068 BP; 41194 A; 30122 C; 32403 G; 39349 T; 0 other;

Query Match 56.0%; Score 1250.8; DB 21; Length 143068;
Best Local Similarity 99.8%; Pred. No. 0;
Matches 1252; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 979 ACCCTTTTCATAGCTCTTGCTGTAGATGGCCCACTCCAAACACAGTGTGTGA 1038
|||||
Db 48253 AGCCTTTTCATAGCTCTTGCTGTAGATGGCCCACTCCAAACACAGTGTGTGA 48312
|||||
Qy 1039 GTCCAGAGTGTAGACACAGGAAGATGCAAGTCACTACACAGGACTCTCGATGT 1098
|||||

Db	48313	GGTCCAGGAGT	GAGACCAGAAAGAAATGGAAGTGA	CTACACAGGACTCCTCGATCGT	48372
QY	1099	CGTGGAAAAG	AAAGTCAATTGGCAGAGCCCTGAAGCCAGTCTTTCAGACAAAGAAGGA	1158	
Db	48373	CGTGGAAAAG	AAAGTCAATTGGCAGAGCCCTGAAGCCAGTCTTTCAGACAAAGAAGGA	48432	
QY	1159	GCCTAGACAG	AAATGACAGATCTCTGCTTTGGRAAATCACACGCTCGCTGCCTTCACAGATG	1218	
Db	48433	GCCTAGACAG	AAATGACAGATCTCTGCTTTGGRAAATCACACGCTCGCTGCCTTCACAGATG	48492	
QY	1219	TGTGATTCAC	AGTGTGAATCTTGCTGTCTACGTTTACCAGGCAGGAAGGCTGAGAGGAG	1278	
Db	48493	TGTGATTCAC	AGTGTGAATCTTGCTGTCTACGTTTACCAGGCAGGAAGGCTGAGAGGAG	48552	
QY	1279	AGACTCCAGC	TGGTGGAAAAACAGTATTTTCCAAACTACCTTCCAGTTCCTCATTTTG	1338	
Db	48553	AGACTCCAGC	TGGTGGAAAAACAGTATTTTCCAAACTACCTTCCAGTTCCTCATTTTG	48612	
QY	1339	AATACAGGC	ATAGAGTTCAGACATTTTAAATAGTAAAAATAAAATTAAGCTGAAAAAC	1398	
Db	48613	AATACAGGC	ATAGAGTTCAGACATTTTAAATAGTAAAAATAAAATTAAGCTGAAAAAC	48672	
QY	1399	TGCAACTTGT	AAATGTGGTAAAGAGTTAGTTTGCAGTTGCTATCATGTCAAAAGTGAATA	1458	
Db	48673	TGCAACTTGT	AAATGTGGTAAAGAGTTAGTTTGCAGTTGCTATCATGTCAAAAGTGAATA	48732	
QY	1459	GCTGTATT	AGTTCACAGAGATAATTCCTAGCTTTGAGCTTAGAATTTTGACGAGTGGTAT	1518	
Db	48733	GCTGTATT	AGTTCACAGAGATAATTCCTAGCTTTGAGCTTAGAATTTTGACGAGTGGTAT	48792	
QY	1519	GTTTGGGAG	ACTGCTGAGTCAACCCAACTAGTTGATTTGGCAGGAGTTGGAAGTGTGTG	1578	
Db	48793	GTTTGGGAG	ACTGCTGAGTCAACCCAACTAGTTGATTTGGCAGGAGTTGGAAGTGTGTG	48852	
QY	1579	ATCTGTGGC	ACATTAGCTTAGTGCAGCAATCTAAGTATGATGCTGTTTGAATCA	1638	
Db	48853	ATCTGTGGC	ACATTAGCTTAGTGCAGCAATCTAAGTATGATGCTGTTTGAATCA	48912	
QY	1639	CAGTATACG	CTCCATCTCAGCTGGATCTCCATCTCTCAGGCTTGCCTGCCA	1698	
Db	48913	CAGTATACG	CTCCATCTCAGCTGGATCTCCATCTCTCAGGCTTGCCTGCCA	48972	
QY	1699	AAAGCCTTT	TGTTTGTATCATTAAGAATCATGCGTTTAAATCACATTCAGT	1758	
Db	48973	AAAGCCTTT	TGTTTGTATCATTAAGAATCATGCGTTTAAATCACATTCAGT	49032	
QY	1759	GTTTCAGT	GCCTTCGAGTCTATGTCATATGTTTCCCTAAATTTGCCAGTGGGAA	1818	
Db	49033	GTTTCAGT	GCCTTCGAGTCTATGTCATATGTTTCCCTAAATTTGCCAGTGGGAA	49092	
QY	1819	CTCCTAAAT	CAAAATGGCTTCTTAATCAAGCTTTTAAACCTATTTGTTAAAGAATGGAAG	1878	
Db	49093	CTCCTAAAT	CAAAATGGCTTCTTAATCAAGCTTTTAAACCTATTTGTTAAAGAATGGAAG	49152	
QY	1879	GTGGAGAAG	CTCCCTGAAGTAAAGCAAGACTTCTCTTACTCGAGCCAACTTAAGAATG	1938	
Db	49153	GTGGAGAAG	CTCCCTGAAGTAAAGCAAGACTTCTCTTACTCGAGCCAACTTAAGAATG	49212	
QY	1939	TTCTTATGT	TGCCAGTGTGTTTCTGATCTGATGCAAGCAAGAACTGGGCTTCTAGA	1998	
Db	49213	TTCTTATGT	TGCCAGTGTGTTTCTGATCTGATGCAAGCAAGAACTGGGCTTCTAGA	49272	
QY	1999	ACCAGGCA	ACTTTGGGAAGTACCTCCCAAGCTGGACTATGGCTCTACCTTCAGGCCACAT	2058	
Db	49273	ACCAGGCA	ACTTTGGGAAGTACCTCCCAAGCTGGACTATGGCTCTACCTTCAGGCCACAT	49332	
QY	2059	GGCTAAAGA	AGGTTTCAGAAAAAGTGGGGCAGACAGCAACTTTTCACTTCATATATTT	2118	
Db	49333	GGCTAAAGA	AGGTTTCAGAAAAAGTGGGGCAGACAGCAACTTTTCACTTCATATATTT	49392	
QY	2119	GTATGATC	CTTAATGAATGCATAAATGTTTAAAGTTGATGGTGAATAATGTAATFACTGTT	2178	
Db	49393	GTATGATC	CTTAATGAATGCATAAATGTTTAAAGTTGATGGTGAATAATGTAATFACTGTT	49452	

QY	2179	TTTAACTATGATTTGGAAAAATAAATGCCTATAACTATGTTGATAAAG	2232
Db	49453	TTTAACTATGATTTGGAAAAATAAATGCCTATAACTATGTTGATAAAG	49506
RESULT 4			
AAA34983			
ID	AAA34983	standard; DNA; 143068 BP.	
XX			
AC	AAA34983;		
XX			
DT	28-JUL-2000	(first entry)	
XX			
DE	Human adenosine receptor related polynucleotide	SEQ ID NO:2672.	
XX			
KW	Human; adenosine receptor; low adenosine antisense oligonucleotide;		
KW	phosphorothioate; impaired respiration; inflammation; allergy;		
KW	allergic disease; bronchoconstriction; inhibitor; antiinflammatory;		
KW	antiallergic; antiasthmatic; cytostatic; analgesic; impaired airway;		
KW	lung disease; ischaemic condition; pulmonary vasoconstriction; asthma;		
KW	respiratory distress syndrome; pain; cystic fibrosis; emphysema;		
KW	pulmonary hypertension; chronic obstructive pulmonary disease; COPD;		
KW	cancer; leukaemia; lymphoma; carcinoma; metastasis; ss.		
XX			
OS	Homo sapiens.		
XX			
PN	WO200009525-A2.		
XX			
PD	24-FEB-2000.		
XX			
PF	03-AUG-1999;	99WO-US17712.	
XX			
PR	03-AUG-1998;	98US-0095212.	
XX			
PA	(UYEC-) UNIV EAST CAROLINA.		
XX			
PI	Nyce JW;		
XX			
DR	WPI; 2000-205971/18.		
XX			
PT	New antisense oligonucleotides useful for treating e.g. pulmonary		
PT	vasoconstriction, inflammation, allergies, asthma, hypertension,		
PT	bronchitis, emphysema, respiratory distress syndrome, ischemia or		
PT	cancers		
XX			
PS	Disclosure; Page 851-882; i343pp; English.		
XX			
CC	The present invention describes a new composition comprising an		
CC	antisense oligonucleotide (ON) with low adenosine (up to 15%), which		
CC	targets nucleic acids involved in bronchoconstriction, allergies, and/or		
CC	inflammation. The ON can have antiinflammatory, antiallergic,		
CC	antiasthmatic, cytostatic and analgesic activities. The compositions are		
CC	useful for the treatment of diseases associated with inflammation,		
CC	impaired airways, including lung disease and diseases whose secondary		
CC	effects afflict the lungs of a subject. They can be used for treating		
CC	e.g. ischaemic conditions, pulmonary vasoconstriction, allergies,		
CC	asthma, impaired respiration, respiratory distress syndrome, pain, cystic		
CC	fibrosis, pulmonary hypertension, emphysema, chronic obstructive		
CC	pulmonary disease (COPD), and cancers which may metastasize to the lungs, including		
CC	carcinomas, and cancers which may metastasize to the lungs, including		
CC	breast and prostate cancer. The reduction of the adenosine content of		
CC	the ONs reduces side effects. The A-containing ONs break down with the		
CC	release of deoxyadenosine which activates adenosine receptors causing the		
CC	bronchoconstriction and inflammation. AAA32313 to AAA35312 represent the		
CC	nucleotide sequences given in the sequence listing from the present		
CC	invention, which correspond to SEQ ID NO:1 to 2815, and then the last		
CC	185 sequences are also called SEQ ID NO:1 to 185, but the sequences		
CC	differ from the previously named sequences. SEQ ID NO:11 to 1680		
CC	(AAA32323 to AAA33992) are specifically claimed ONs from the present		
CC	invention. N.B. Sequences given in the disclosure of the present		
CC	invention do not match up with their corresponding SEQ ID NO: sequences		
CC	given in the sequence listing.		

carcinomas, and cancers which may metastasize to the lungs, including breast and prostate cancer. The reduction of the adenosine content of the ONS reduces side effects. The A-containing ONS break down with the release of deoxyadenosine which activates adenosine receptors causing bronchoconstriction and inflammation. AAA32313 to AAA35312 represent the nucleotide sequences given in the sequence listing from the present invention, which correspond to SEQ ID NO:1 to 185, and then the last 185 sequences are also called SEQ ID NO:1 to 185, but the sequences differ from the previously named sequences. SEQ ID NO:11 to 1680 (AAA32323 to AAA33992) are specifically claimed ONS from the present invention. N.B. Sequences given in the disclosure of the present invention do not match up with their corresponding SEQ ID NO: sequences given in the sequence listing.

Sequence 143068 BP; 41194 A; 30126 C; 32402 G; 39346 T; 0 other;

Query Match 56.0%; Score 1250.8; DB 21; Length 143068;
Best Local Similarity 99.8%; Pred. No. 0;
Matches 1252; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY	979	AGCCCTTTTACATAGCTCTGGCTGTAGGATTGCCACTCCAAAACACAGTGTGGA	1038
DB	48253	AGCCCTTTTACATAGCTCTGGCTGTAGGATTGCCACTCCAAAACACAGTGTGGA	48312
QY	1039	GGTCCAGAGTGAGACCGAAGAATGTGAAGTCACTACACAGGACTCCTCGATGTT	1098
DB	48313	GGTCCAGAGTGAGACCGAAGAATGTGAAGTCACTACACAGGACTCCTCGATGTT	48372
QY	1099	CGTGGAAAAGGAAGTCAATTTGGCAGAGCCCTTGAAGCAGTCTTCAGGACAAAGGA	1158
DB	48373	CGTGGAAAAGGAAGTCAATTTGGCAGAGCCCTTGAAGCAGTCTTCAGGACAAAGGA	48432
QY	1159	GCCTAGAGACAGAAATGACAGATCTCTGCTTTGGAATACACAGTCTGGCTTCACAGATG	1218
DB	48433	GCCTAGAGACAGAAATGACAGATCTCTGCTTTGGAATACACAGTCTGGCTTCACAGATG	48492
QY	1219	TGTGATTCACAGTCAATCTGGTGTCTACGTTACAGCAGGAGAGGCTGAGAGGAGAG	1278
DB	48493	TGTGATTCACAGTCAATCTGGTGTCTACGTTACAGCAGGAGAGGCTGAGAGGAGAG	48552
QY	1279	AGACTCCAGCTGGTGTGGAACAGTATTTCCAAACTACCTTCCAGTCTCCTCATTTTG	1338
DB	48553	AGACTCCAGCTGGTGTGGAACAGTATTTCCAAACTACCTTCCAGTCTCCTCATTTTG	48612
QY	1339	AATACAGGATAGAGTTCAGACTTTTAAATAGTAAATATAAATTAAGCTGAAAC	1398
DB	48613	AATACAGGATAGAGTTCAGACTTTTAAATAGTAAATATAAATTAAGCTGAAAC	48672
QY	1399	TGCAACTTGTAAATGTGTAAGAGTGTAGTTTGTAGTTGCTATCATGTCAACAGTGAAT	1458
DB	48673	TGCAACTTGTAAATGTGTAAGAGTGTAGTTTGTAGTTGCTATCATGTCAACAGTGAAT	48732
QY	1459	GCTGTATTAGTACAGAGATAATTTAGCTTTGAGTTGAGTTAGAAATTTTGACGAGTGTAT	1518
DB	48733	GCTGTATTAGTACAGAGATAATTTAGCTTTGAGTTGAGTTAGAAATTTTGACGAGTGTAT	48792
QY	1519	GTITGGGAGACTGCTGAGTCAACCAATAGTTTGTAGTTGCGCAGGAGTTGGAAGTGTG	1578
DB	48793	GTITGGGAGACTGCTGAGTCAACCAATAGTTTGTAGTTGCGCAGGAGTTGGAAGTGTG	48852
QY	1579	ATCTGTGGGACATAGCTATGTGATGACATCATCAATCAATGATGCTGTTGAATCA	1638
DB	48853	ATCTGTGGGACATAGCTATGTGATGACATCATCAATCAATGATGCTGTTGAATCA	48912
QY	1639	CAGTATAGCTTCCATCGCTGTCATCTAGCTGGATCTCCATCTCTCAGGCTTCTGCGCA	1698
DB	48913	CAGTATAGCTTCCATCGCTGTCATCTAGCTGGATCTCCATCTCTCAGGCTTCTGCGCA	48972
QY	1699	AAAGCCTTTTGTGTTTGTGATCATTTAAGTCAATGAGTCAATGATGCTGTTGAGT	1758
DB	48973	AAAGCCTTTTGTGTTTGTGATCATTTAAGTCAATGAGTCAATGATGCTGTTGAGT	49032
QY	1759	GTITTCAGTGTCTCGCAGATGCTTGTGATGCTCATATTTGCTTAATTTGCGAGTGGAA	1818

DB	49033	GTITTCAGTGTCTCGCAGATGCTTGTGATGCTCATATTTGCTTATTTGCCAGTGGAA	49092
QY	1819	CTCCTAAATCAAAATTTGGCTTCTAAATCAAAAGCTTTTAAACCCCTATTGGTAAAGTGAAG	1878
DB	49093	CTCCTAAATCAAAATTTGGCTTCTAAATCAAAAGCTTTTAAACCCCTATTGGTAAAGTGAAG	49152
QY	1879	GTGGAGAGCTCCCTGAACTAAGCAAGACTTTTCCCTTTAGTCGAGCCAAAGTAAAGATG	1938
DB	49153	GTGGAGAGCTCCCTGAACTAAGCAAGACTTTTCCCTTTAGTCGAGCCAAAGTAAAGATG	49212
QY	1939	TTCTTATGTTGCCAGTGTGTTTCTGATCTGATGCAAGCAAGAAACACACCTGCGGCTTCTAGA	1998
DB	49213	TTCTTATGTTGCCAGTGTGTTTCTGATCTGATGCAAGCAAGAAACACACCTGCGGCTTCTAGA	49272
QY	1999	ACCAGCAACTTTGGGAACTAGACTCCCAAGCTGGACTATGGCTCTACTTTTCAGGCCACAT	2058
DB	49273	ACCAGCAACTTTGGGAACTAGACTCCCAAGCTGGACTATGGCTCTACTTTTCAGGCCACAT	49332
QY	2059	GGCTAAAGAAAGTTTTCAGAAAGAAAGTGGGACAGACAGCAAACTTTCCACCTTCATATATTT	2118
DB	49333	GGCTAAAGAAAGTTTTCAGAAAGAAAGTGGGACAGACAGCAAACTTTCCACCTTCATATATTT	49392
QY	2119	GTATGATCTCTAAATGAATGCAATAAAATGTTAAGTTGATGGTGAATGTAATGTAATGCTTT	2178
DB	49393	GTATGATCTCTAAATGAATGCAATAAAATGTTAAGTTGATGGTGAATGTAATGTAATGCTTT	49452
QY	2179	TTTTAAACACTATCATTTGGAAATAAATCAATCAATGCTATAACTATGTTGATAAAAG	2232
DB	49453	TTTTAAACACTATCATTTGGAAATAAATCAATCAATGCTATAACTATGTTGATAAAAG	49506

RESULT 6
ABL68124
ID ABL68124 standard; DNA; 143068 BP.
XX ABL68124;
XX AC
XX XX
DT 15-MAY-2002 (first entry)
XX Ovary cancer related gene sequence SEQ ID NO:6461.
DE Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
KW stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
KW cytostatic; gene therapy; antineoplastic; Wilms' tumour; adenocarcinoma;
KW gene; ds.
XX Homo sapiens.
OS WO200194629-A2.
PN 13-DEC-2001.
XX 30-MAY-2001; 2001WO-US10838.
XX 05-JUN-2000; 2000US-209473P.
PR 05-JUN-2000; 2000US-209531P.
PR 18-SEP-2000; 2000US-233133P.
PR 18-SEP-2000; 2000US-233617P.
PR 20-SEP-2000; 2000US-234009P.
PR 20-SEP-2000; 2000US-234034P.
PR 20-SEP-2000; 2000US-234052P.
PR 22-SEP-2000; 2000US-234509P.
PR 22-SEP-2000; 2000US-234567P.
PR 25-SEP-2000; 2000US-234923P.
PR 25-SEP-2000; 2000US-234924P.
PR 25-SEP-2000; 2000US-235077P.
PR 25-SEP-2000; 2000US-235082P.
PR 25-SEP-2000; 2000US-235134P.
PR 25-SEP-2000; 2000US-235280P.
PR 25-SEP-2000; 2000US-235637P.
PR 26-SEP-2000; 2000US-235638P.
PR 27-SEP-2000; 2000US-235711P.

RESULT 7

AAA35151
ID AAA35151 standard; DNA; 149412 BP.

XX AC AAA35151;

XX DT 28-JUL-2000 (first entry)

XX DE Human adenosine receptor related polynucleotide 2nd SEQ ID NO:25.

XX KW Human; adenosine receptor; low adenosine antisense oligonucleotide; phosphorothioate; impaired respiration; inflammation; allergy;
KW allergic disease; bronchoconstriction; inhibitor; antiinflammatory;
KW antiallergic; antiallergic; cytotatic; analgesic; impaired airway;
KW lung disease; ischaemic condition; pulmonary vasoconstriction; asthma;
KW respiratory distress syndrome; pain; cystic fibrosis; emphysema;
KW pulmonary hypertension; chronic obstructive pulmonary disease; COPD;
KW cancer; leukaemia; lymphoma; carcinoma; metastasis; ss.

XX OS Homo sapiens.

XX PN WO200009525-A2.

XX PD 24-FEB-2000.

XX PF 03-AUG-1999; 99WO-US17712.

XX PR 03-AUG-1998; 98US-0095212.

XX PA (UYEC-) UNIV EAST CAROLINA.

XX PI Nyce JW;

XX DR WPI; 2000-205971/18.

XX PT New antisense oligonucleotides useful for treating e.g. pulmonary
PT vasoconstriction, inflammation, allergies, asthma, hypertension,
PT bronchitis, emphysema, respiratory distress syndrome, ischemia or
PT cancers -

XX PS Disclosure; Page 1138-1171; 1343pp; English.

XX CC The present invention describes a new composition comprising an
XX antisense oligonucleotide (ON) with low adenosine (up to 15%), which
XX targets nucleic acids involved in bronchoconstriction, allergies, and/or
XX inflammation. The ON can have antiinflammatory, antiallergic,
XX antiasthmatic, cytotatic and analgesic activities. The compositions are
XX useful for the treatment of diseases associated with inflammation,
XX impaired airways, including lung disease and diseases whose secondary
XX effects afflict the lungs of a subject. They can be used for treating
XX e.g. ischaemic conditions, pulmonary vasoconstriction, allergies, cystic
XX asthma, impaired respiration, respiratory distress syndrome, pain, chronic
XX fibrosis, pulmonary hypertension, emphysema, chronic obstructive
XX pulmonary disease (COPD), and cancers such as leukaemias, lymphomas,
XX carcinomas, and cancers which may metastasize to the lungs, including
XX breast and prostate cancer. The reduction of the adenosine content of
XX the ONs reduces side effects. The A-containing ONs break down with the
XX release of deoxyadenosine which activates adenosine receptors causing
XX bronchoconstriction and inflammation. AAA3313 to AAA35312 represent the
XX nucleotide sequences given in the sequence listing from the present
XX invention, which correspond to SEQ ID NO:1 to 2815, and then the last
XX 185 sequences are also called SEQ ID NO:1 to 185, but the sequences
XX differ from the previously named sequences. SEQ ID NO:11 to 1680
XX (AAA3232 to AAA3392) are specifically claimed ONs from the present
XX invention. N.B. Sequences given in the disclosure of the present
XX invention do not match up with their corresponding SEQ ID NO: sequences
XX given in the sequence listing.

XX SQ Sequence 149412 BP; 43049 A; 31388 C; 33852 G; 41123 T; 0 other;

Query Match 56.0%; Score 1250.8; DB 21; Length 149412;
Best Local Similarity 99.8%; Pred. No. 0;

	Matches	1252;	Conservative	0;	Mismatches	2;	Indels	0;	Gaps	0;
QY	979	AGCCTTTTTCACATAGCTCTTGGCTGTAGGATTGCCCACTCCAAAACACAGTGTGTGGA	1038							
DB	54597	AGCCTTTTTCACATAGCTCTTGGCTGTAGGATTGCCCACTCCAAAACACAGTGTGTGGA	54656							
QY	1039	GGTCCAGGAGTGAGACAGGAAAGATGTGAAAGTGTACTACAAAGGACTCTCCGATGGT	1098							
DB	54657	GGTCCAGGAGTGAGACAGGAAAGATGTGAAAGTGTACTACAAAGGACTCTCCGATGGT	54716							
QY	1099	CGTGGAAAAGAAAGTCAATTTGGCAGAGCCCTGAAAGCCAGTCTTTCAGGACAAAGAGGA	1158							
DB	54717	CGTGGAAAAGAAAGTCAATTTGGCAGAGCCCTGAAAGCCAGTCTTTCAGGACAAAGAGGA	54776							
QY	1159	GCTTAGACAGAGAAATGACAGATCTCTGCTTTGGAAATACACGCTCTGGCTTTCACAGATG	1218							
DB	54777	GCTTAGACAGAGAAATGACAGATCTCTGCTTTGGAAATACACGCTCTGGCTTTCACAGATG	54836							
QY	1219	TGTGATTTCAGAGTGTGAATCTTGGTGTCTAGCTTACCAGGAGGAGGCTGAGAGGAGAG	1278							
DB	54837	TGTGATTTCAGAGTGTGAATCTTGGTGTCTAGCTTACCAGGAGGAGGCTGAGAGGAGAG	54896							
QY	1279	AGACTCCAGCTGGGTTGGAAAACAGATATTTTCCAAACTACCTTCCAGTTCCTCATTTTGG	1338							
DB	54897	AGACTCCAGCTGGGTTGGAAAACAGATATTTTCCAAACTACCTTCCAGTTCCTCATTTTGG	54956							
QY	1339	AATACAGGCATAGAGTTTCAGACTTTTTTAAATAGTAAAAATAAAATTAAGAGCTGAAAC	1398							
DB	54957	AATACAGGCATAGAGTTTCAGACTTTTTTAAATAGTAAAAATAAAATTAAGAGCTGAAAC	55016							
QY	1399	TGCAACTTGTAAATGTGTTAAAGAGTTAGTTTGGATTGCTATCATGTCAACACGTGAAAT	1458							
DB	55017	TGCAACTTGTAAATGTGTTAAAGAGTTAGTTTGGATTGCTATCATGTCAACACGTGAAAT	55076							
QY	1459	GCTGTATTAGTCACAGAGATTAATTTCTAGCTTTGAGCTTAAGAAATTTTCAGCAGGTGGTAT	1518							
DB	55077	GCTGTATTAGTCACAGAGATTAATTTCTAGCTTTGAGCTTAAGAAATTTTCAGCAGGTGGTAT	55136							
QY	1519	GTTTGGGAGACTGCTGAGTCAACCAATAGTTTGTGATTGGCAGGAGTTGGAAAGTGTGTG	1578							
DB	55137	GTTTGGGAGACTGCTGAGTCAACCAATAGTTTGTGATTGGCAGGAGTTGGAAAGTGTGTG	55196							
QY	1579	ATCTGTGGGCACATAGGCTATGTGCATGCAGCATCTAAGTAATGATGCTGTTGAATCA	1638							
DB	55197	ATCTGTGGGCACATAGGCTATGTGCATGCAGCATCTAAGTAATGATGCTGTTGAATCA	55256							
QY	1639	CAGTATACGCTCCATCGCTGCTCATCTCAGCTGGATCTCCATTTCTCTCAGGCTTGTGCCA	1698							
DB	55257	CAGTATACGCTCCATCGCTGCTCATCTCAGCTGGATCTCCATTTCTCTCAGGCTTGTGCCA	55316							
QY	1699	AAAGCCTTTTGTGTTTGTGTTTGTATCATATTGAAGTCATGCGTTTAAATCATCTCGAGT	1758							
DB	55317	AAAGCCTTTTGTGTTTGTGTTTGTATCATATTGAAGTCATGCGTTTAAATCATCTCGAGT	55376							
QY	1759	GTTTCAGTGTTCGAGATGCTCTGATCTCATATTGTTCCCTAATTTGCCAGTGGGAA	1818							
DB	55377	GTTTCAGTGTTCGAGATGCTCTGATCTCATATTGTTCCCTAATTTGCCAGTGGGAA	55436							
QY	1819	CTCCTAAATCAAAATTTGGCTTCTTAATCAAAAGCTTTTAAACCCCTATTGGTAAAGAAATG	1878							
DB	55437	CTCCTAAATCAAAATTTGGCTTCTTAATCAAAAGCTTTTAAACCCCTATTGGTAAAGAAATG	55496							
QY	1879	GTGGAGAGCTCCCTGGAAGTAAGAAAGACTTTTCTCTTCTAGTCGAGCCCAAGTAAAGATG	1938							
DB	55497	GTGGAGAGCTCCCTGGAAGTAAGAAAGACTTTTCTCTTCTAGTCGAGCCCAAGTAAAGATG	55556							
QY	1939	TTCTTATGTTGCCAGTGTGTTTCTGATCTGATGAAGCAAGAAACACTGGGCTTCTAGA	1998							
DB	55557	TTCTTATGTTGCCAGTGTGTTTCTGATCTGATGAAGCAAGAAACACTGGGCTTCTAGA	55616							
QY	1999	ACCAGCAACTTGGGAACTAGACTCCCAAGCTGAGCTATGGCTCTACTTTTCAGGCCACAT	2058							
DB	55617	ACCAGCAACTTGGGAACTAGACTCCCAAGCTGAGCTATGGCTCTACTTTTCAGGCCACAT	55676							

CC family, degenerate oligo primers were designed corresp. to the conserved sequences R79167 in the second and R79168 in the third transmembrane domains of the MIP-1alpha/RANTES receptor, the IL-8 receptors and the HUMSTRS orphan receptor (GenBank Accession #M99293. The degenerate oligo incorporating EcoRI and XhoI sites at their 5' ends are Q96299 and Q96300. Amplification of cDNA derived from MM6 cells with the primers yielded a number of PCR products. One cDNA appeared to encode a novel protein. To obtain a full-length version of this clone, a MM6 cDNA library was constructed in pPROG and probed with the PCR product. A 2.1 kb cDNA clone was obtained. Analysis of additional clones in the MM6 cDNA library revealed a second sequence that was identical to the 2.1 kb cDNA sequence first obtained. The 5' UTR through the putative seventh transmembrane domain but contained a different cytoplasmic tail. The second sequence appears to represent alternative splicing of the carboxyl-terminal tail of the MCP-1R protein. The two sequences are denoted MCP-1RA and MCP-1RB (see Q96297/R79165 & Q96298/R79166). Active mature MCP-1RA has a predicted mol. wt. of about 42,000 daltons. MCP-1RB has a mol. wt. of about 41,000 daltons.

XX
SQ Sequence 1979 BP; 530 A; 434 C; 452 G; 563 T; 0 other;

Query Match 43.9%; Score 980; DB 16; Length 1979;
Best Local Similarity 100.0%; Pred. No. 9,8e-263;
Matches 980; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGATTGAACAAGACGATTTCCCAAGTATATCCCAACATGCTGCCACATCTGTTCT 60
DB 42 GGATTGAACAAGACGATTTCCCAAGTATATCCCAACATGCTGCCACATCTGTTCT 101
QY 61 CGGTTTATCAGAAATACCAACGAGAGCGGTGAAGAGTACACACCTTTTGTATTATGAT 120
DB 102 CGGTTTATCAGAAATACCAACGAGAGCGGTGAAGAGTACACACCTTTTGTATTATGAT 161
QY 121 TACGGTCTCCCTGTCATATAATTTGACGTGAAGCAAAATTTGGGCCCCAACCTCTGCCCTCCG 180
DB 162 TACGGTCTCCCTGTCATATAATTTGACGTGAAGCAAAATTTGGGCCCCAACCTCTGCCCTCCG 221
QY 181 CTCTACTCGCTGGTGTTCATCTTTGGTTTGTGGGCAACATGCTGGTGGTCTCTATCTTA 240
DB 222 CTCTACTCGCTGGTGTTCATCTTTGGTTTGTGGGCAACATGCTGGTGGTCTCTATCTTA 281
QY 241 ATAACTGCAAAAAGCTGAAGTGTCTGACGTGACATTTACCTGCTCAACCTGCCATCTCT 300
DB 282 ATAACTGCAAAAAGCTGAAGTGTCTGACGTGACATTTACCTGCTCAACCTGCCATCTCT 341
QY 301 GATCTGCTTTTCTTATTACTCTCCCATTTGCTGGGCTCACTGCTGCTCAAAATGAGTGGTC 360
DB 342 GATCTGCTTTTCTTATTACTCTCCCATTTGCTGGGCTCACTGCTGCTCAAAATGAGTGGTC 401
QY 361 TTTGGGAATGCAATGTGCAAAATTTACAGGGGTGTATACATCGGTATTATTTGGCGGA 420
DB 402 TTTGGGAATGCAATGTGCAAAATTTACAGGGGTGTATACATCGGTATTATTTGGCGGA 461
QY 421 ATCTTCTTCATCATCTCCCTGACAAATGATAGATACCTGGCTATTGTCCATGCTGTGTT 480
DB 462 ATCTTCTTCATCATCTCCCTGACAAATGATAGATACCTGGCTATTGTCCATGCTGTGTT 521
QY 481 GCTTTAAAAGCCAGGAGCGGTCAACCTTTGGGGTGTGTGACAAAGTGTATGATCGGTGGTG 540
DB 522 GCTTTAAAAGCCAGGAGCGGTCAACCTTTGGGGTGTGTGACAAAGTGTATGATCGGTGGTG 581
QY 541 GCTGTGTTTGTCTGTGCTCCAGGAATCATCTTTACTAATGCCAGAAAGAGATTCTGTT 600
DB 582 GCTGTGTTTGTCTGTGCTCCAGGAATCATCTTTACTAATGCCAGAAAGAGATTCTGTT 641
QY 601 TATGCTCTGCGCCCTTATTTTCCAGGAGATGGAATAATTTCCACACATAATAGGAAAC 660
DB 642 TATGCTCTGCGCCCTTATTTTCCAGGAGATGGAATAATTTCCACACATAATAGGAAAC 701
QY 661 ATTTTGGGCTGGTCTCCCGCTGCTCATATGATGCTGCTACTCGGGAATCTGAAA 720
DB 702 ATTTTGGGCTGGTCTCCCGCTGCTCATATGATGCTGCTACTCGGGAATCTGAAA 761

QY 1759 GTTTCAGTGTTCGAGATGTCCTTGATGCTCATATTTGTTCCCTAAATTTGCCAGTGGAA 1818
DB 5537 GTTTCAGTGTTCGAGATGTCCTTGATGCTCATATTTGTTCCCTAAATTTGCCAGTGGAA 55436
QY 1819 CTCCTAAATCAAAATGGCTTCTAATCAAGCTTTTAAACCTATTTGGTAAAGATGGAAG 1878
DB 55437 CTCCTAAATCAAAATGGCTTCTAATCAAGCTTTTAAACCTATTTGGTAAAGATGGAAG 55496
QY 1879 GTGAGAGCTCCCTGAAGTAAAGCAAGACTTTCTCTTAGTCGAGCAAGTTAAAGATG 1938
DB 55497 GTGAGAGCTCCCTGAAGTAAAGCAAGACTTTCTCTTAGTCGAGCAAGTTAAAGATG 55556
QY 1939 TTTCTATTTGCCAGTGTGTTCTGATCTGATGCAAGCAAGACACTGGCTTCTAGA 1998
DB 55557 TTTCTATTTGCCAGTGTGTTCTGATCTGATGCAAGCAAGACACTGGCTTCTAGA 55616
QY 1999 ACCAGGCAACTTGGGAAGTACTGCCAAGCTGGACTATGGCTCTACCTTTTTCAGGCGACAT 2058
DB 55617 ACCAGGCAACTTGGGAAGTACTGCCAAGCTGGACTATGGCTCTACCTTTTTCAGGCGACAT 55676
QY 2059 GCCTAAAGAAAGTTTCAGAAAGAGTGGGACAGAGCAAGACTTTCACCTTCATATATT 2118
DB 55677 GCCTAAAGAAAGTTTCAGAAAGAGTGGGACAGAGCAAGACTTTCACCTTCATATATT 55736
QY 2119 GTATGATCTTAATGATGATCAATAAATGTTAAGTTGATGATGATGATGATGATGATGAT 2178
DB 55737 GTATGATCTTAATGATGATCAATAAATGTTAAGTTGATGATGATGATGATGATGATGAT 55796
QY 2179 TTTTAAACACTATGATTTGGAAATAAATCAATGCTATTAATGATGATGATGATGATGATGAT 2232
DB 55797 TTTTAAACACTATGATTTGGAAATAAATCAATGCTATTAATGATGATGATGATGATGATGAT 55850
RESULT 9
AAQ96298
ID AAQ96298 standard; cDNA; 1979 BP.
XX AAQ96298;
AC AAQ96298;
XX
DT 29-DEC-1995 (first entry)
DE Human monocyte chemoattractant protein-1 receptor MCP-1RB.
XX
KW Monocyte chemoattractant protein-1 receptor; MCP-1R; chemokine; ss.
XX
OS Homo sapiens.
XX
FH Key
FT CDS
FT Location/Qualifiers
81..1160
/*tag= a
W09519436-A.
XX
XX 20-JUL-1995.
XX
XX 11-JAN-1995; 95WO-US00476.
XX
XX 13-JAN-1994; 94US-0182962.
XX
XX (REGC) UNIV CALIFORNIA.
XX
XX Charo I, Coughlin S;
XX
XX WPI; 1995-263866/34.
XX
XX P-PSDB; AAR79166.
XX
XX DNA encoding monocyte chemo-attractant protein-1 receptor - used partic.
XX
XX for identifying antagonists and for treating diseases characterised by
XX
XX monocyte infiltrates
XX
XX Disclosure; Fig 2; 84pp; English.
XX
XX To identify and clone new members of the chemokine receptor gene
XX
XX

QY 721 ACCCTGCTTCGGTGTGGAACGAGAGGAGGATAGGCGAGTGAGAGTCACTTTCCACC 780
|||||
Db 762 ACCCTGCTTCGGTGTGGAACGAGAGGAGGATAGGCGAGTGAGAGTCACTTTCCACC 821
|||||
QY 781 ATCATGATGTTTACTTCTCTGGAAGTCCCTATATACATGTCATCTCTGGAACACC 840
|||||
Db 822 ATCATGATGTTTACTTCTCTGGAAGTCCCTATATACATGTCATCTCTGGAACACC 881
|||||
QY 841 TTCAGGAATTCCTGCGCTGAGTAAGTGTGAAAGCACCAGTCAACTGGACCCAGCCACG 900
-Db 882 TTCAGGAATTCCTGCGCTGAGTAAGTGTGAAAGCACCAGTCAACTGGACCCAGCCACG 941
|||||
QY 901 CAGGTGACAGAGACTCTTGGGATGACTACCTGCTGCATCAATCCCATCATCTATGCTTC 960
Db 942 CAGGTGACAGAGACTCTTGGGATGACTACCTGCTGCATCAATCCCATCATCTATGCTTC 1001
QY 961 GTTGGGAGAGGTTCAAGAG 980
|||||
Db 1002 GTTGGGAGAGGTTCAAGAG 1021

RESULT 10

AAS12140

ID AAS12140 standard; DNA; 1083 BP.

XX

AC AAS12140;

XX

DT 04-DEC-2001 (first entry)

XX

DE Human wild-type CCR2-64V polynucleotide.

XX

KW Human; CCR2 receptor; CCR2-64I; CCR2-64V; gene therapy; atherosclerosis;
KW single nucleotide polymorphism; hypercholesterolaemia; ds.

XX

OS Homo sapiens.

XX

PN WO200162796-A1.

XX

PD 30-AUG-2001.

XX

PF 22-FEB-2001; 2001WO-GB00755.

XX

PR 22-FEB-2000; 2000GB-0004183.

XX

PA (SMIK) SMITHKLINE BEECHAM PLC.

XX

PI Valdes AM, Groot PHE, Spurr NK;

XX

DR WPI; 2001-550086/61.

XX

DR P-PSDB; AAU07614.

XX

Diagnosing atherosclerosis or susceptibility to atherosclerosis in a
subject, by determining a single nucleotide polymorphism in specific
codon of a polynucleotide encoding human CCR2 receptor in genome of the
subject -

XX

Claim 3; Page 20-21; 28pp; English.

XX

The invention relates to diagnosing atherosclerosis (or susceptibility
to) in a subject by determining expression or activity of the human
CCR2-64I polypeptide (a polymorphic variant form of the human
CCR2 receptor) or the CCR2-64V polypeptide (human CCR2 receptor), by screening
for a single nucleotide polymorphism in codon 64 of the polynucleotide
encoding the CCR2 receptor. This results in production of CCR2-64I,
whereby polymorphic variants are associated with a lower incidence of
atherosclerosis. The presence or amount of CCR2-64I/V in a sample can
also be analysed. The sequences of the invention can be used for
predicting the response of a patient to drug treatment, for predicting
the disease outcome in a patient and also for the production of a
treatment for hypercholesterolaemia. The sequence represents DNA encoding
the wild-type receptor polypeptide CCR2-64V.

XX

SQ Sequence 1083 BP; 255 A; 260 C; 247 G; 321 T; 0 other;

Query Match 42.2%; Score 941; DB 22; Length 1083;

Best Local Similarity 100.0%; Pred. No. 5.5e-252;

Matches 941; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 40 ATGCTCTCCACATCTCGTCTCGGTTTATCAGAAATACCAACGAGAGCGGTGAAGAGTC 99

Db 1 ATGCTCTCCACATCTCGTCTCGGTTTATCAGAAATACCAACGAGAGCGGTGAAGAGTC 60

QY 100 ACCACCTTTTGTATGATTACGGTGTCCCTGTATATAATTTGACGTGAACCAATTT 159

Db 61 ACCACCTTTTGTATGATTACGGTGTCCCTGTATATAATTTGACGTGAACCAATTT 120

QY 160 GGGGCCCCAACTCCTGCTCCGCTCTACTCGCTGGTGTCTATCTTTGTTGGGCAAC 219

Db 121 GGGGCCCCAACTCCTGCTCCGCTCTACTCGCTGGTGTCTATCTTTGTTGGGCAAC 180

QY 220 ATGCTGGTGTCTCATCTTAATAAATCCAAAAGCTGAAGTCTTGACTGACATTTAC 279

Db 181 ATGCTGGTGTCTCATCTTAATAAATCCAAAAGCTGAAGTCTTGACTGACATTTAC 240

QY 280 CTGCTCAACCTGGCCATCTCTGATCTGCTTTTCTTATTACTCTCCCATTTGGGCTCAC 339

Db 241 CTGCTCAACCTGGCCATCTCTGATCTGCTTTTCTTATTACTCTCCCATTTGGGCTCAC 300

QY 340 TCTGCTGCAATGAGTGGTCTTTGGGAATGCAATGTCAAAATTTATTCACAGGCTGTAT 399

Db 301 TCTGCTGCAATGAGTGGTCTTTGGGAATGCAATGTCAAAATTTATTCACAGGCTGTAT 360

QY 400 CACATCGGTTATTTTGGGGAATCTTCTTCATCATCTCTGACATTCGATACATACCTG 459

Db 361 CACATCGGTTATTTTGGGGAATCTTCTTCATCATCTCTGACATTCGATACATACCTG 420

QY 460 GCTATTGTCATGCTGCTTTGCTTTTAAAGCCAGGACGGTCACTTTGGGCTGTGACA 519

Db 421 GCTATTGTCATGCTGCTTTGCTTTTAAAGCCAGGACGGTCACTTTGGGCTGTGACA 480

QY 520 AGTGTGATCACTGGTGTGGTGTGCTTCTGCTCCAGGATCATCTTTACTATAA 579

Db 481 AGTGTGATCACTGGTGTGGTGTGCTTCTGCTCCAGGATCATCTTTACTATAA 540

QY 580 TGCAGAAAGAGATCTGTTTATGCTGTGGCCCTTATTTTCCAGGAGATGGAATAAT 639

Db 541 TGCAGAAAGAGATCTGTTTATGCTGTGGCCCTTATTTTCCAGGAGATGGAATAAT 600

QY 640 TTCCACACAATATGAGGACATTTTGGGCTGTGCTCCGCTGCTCATATGTCATC 699

Db 601 TTCCACACAATATGAGGACATTTTGGGCTGTGCTCCGCTGCTCATATGTCATC 660

QY 700 TGCTACTCGGGAATCTTGAACCCCTGCTTCGGTGTGCGAAAGAGAGGATAGG 759

Db 661 TGCTACTCGGGAATCTTGAACCCCTGCTTCGGTGTGCGAAAGAGAGGATAGG 720

QY 760 GCAGTGAGAGTCACTTCCACCATCATGATGTTTCTTCTTGAGGATCCCTTATAAC 819

Db 721 GCAGTGAGAGTCACTTCCACCATCATGATGTTTCTTCTTGAGGATCCCTTATAAC 780

QY 820 ATTTGTCATCTCTGGAACCTTCCAGGAATCTTCCGCTGAGTAACTGTGAAGACACC 879

Db 781 ATTTGTCATCTCTGGAACCTTCCAGGAATCTTCCGCTGAGTAACTGTGAAGACACC 840

QY 880 AGTCAACTGGACCAAGCCAGGAGTGTGAGAGACTCTTGGGATGACTCACTGTGTCATC 939

Db 841 AGTCAACTGGACCAAGCCAGGAGTGTGAGAGACTCTTGGGATGACTCACTGTGTCATC 900

QY 940 AATCCCATCATCTATGCTTCGTTGGGAGAGTTTCAAGAG 980

Db 901 AATCCCATCATCTATGCTTCGTTGGGAGAGTTTCAAGAG 941

RESULT 11

AAS12139

	AAS12139 standard; DNA; 1083 BP.	
XX	AAS12139;	
XX	04-DEC-2001 (first entry)	
XX	Human CCR2-64I polymorphic variant polynucleotide.	
XX	Human; CCR2 receptor; CCR2-64I; CCR2-64V; gene therapy; atherosclerosis;	
XX	single nucleotide polymorphism; hypercholesterolaemia; ds.	
XX	Homo sapiens.	
XX	Key Location/Qualifiers	
XX	variation replace(190,G)	
XX	/tag= a	
XX	/standard_name= "Single nucleotide polymorphism"	
XX	WO200162796-A1.	
XX	30-AUG-2001.	
XX	22-FEB-2001; 2001WO-GB00755.	
XX	22-FEB-2000; 2000GB-0004183.	
XX	(SMIK) SMITHLINE BEECHAM PLC.	
XX	Valdes AM, Groot PHE, Spurr NK;	
XX	WPI; 2001-550086/61.	
XX	P-PADB; AAU07613.	
XX	Diagnosing atherosclerosis or susceptibility to atherosclerosis in a	
XX	subject, by determining a single nucleotide polymorphism in specific	
XX	codon of a polynucleotide encoding human CCR2 receptor in genome of the	
XX	subject -	
XX	Claim 3; Page 20; 28pp; English.	
XX	The invention relates to diagnosing atherosclerosis (or susceptibility	
XX	to) in a subject by determining expression or activity of the human	
XX	CCR2-64I polypeptide (a polymorphic variant form of the human CCR2	
XX	receptor) or the CCR2-64V polypeptide (human CCR2 receptor), by screening	
XX	for a single nucleotide polymorphism in codon 64 of the polynucleotide	
XX	encoding the CCR2 receptor. This results in production of CCR2-64I,	
XX	whereby polymorphic variants are associated with a lower incidence of	
XX	atherosclerosis. The presence or amount of CCR2-64I/V in a sample can	
XX	also be analysed. The sequences of the invention can be used for	
XX	predicting the response of a patient to drug treatment, for predicting	
XX	the disease outcome in a patient and also for the production of a	
XX	treatment for hypercholesterolaemia. The sequence represents DNA encoding	
XX	the polymorphic variant polypeptide CCR2-64I.	
XX	Sequence 1083 BP; 256 A; 260 C; 246 G; 321 T; 0 other;	
XX	Query Match 42.1%; Score 939.4; DB 22; Length 1083;	
XX	Best Local Similarity 99.9%; Pred. No. 1.5e-251;	
XX	Matches 940; Conservative 0; Mismatches 1; Indels 0; Gaps 0;	
QY	40 ATGCTGTCCACATCTCGTTCGGTTTATCAGAAATACCACGAGAGCGGTGAAGAAGTC 99	
DB	1 ATGCTGTCCACATCTCGTTCGGTTTATCAGAAATACCACGAGAGCGGTGAAGAAGTC 60	
QY	100 ACCACTTTTGGATTGATGATACGGTGCTCCCTGCATATAAATTTGACGTGAAGCAAAAT 159	
DB	61 ACCACTTTTGGATTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 120	
QY	160 GGGGCCCAACTCCTGCCTCCGCTACTACTCGTGGTGTTCATCTTTGGTTTGGGGCAAC 219	
DB	121 GGGGCCCAACTCCTGCCTCCGCTACTACTCGTGGTGTTCATCTTTGGTTTGGGGCAAC 180	

[illegible]

XX	Human immune system associated gene SEQ ID NO: 308.
XX	
XX	Human; immune system disease; cytosine methylation; antiasthmatic;
KW	antiarteriosclerotic; antianemic; cytostatic; nootropic;
KW	neuroprotective; anti-HIV; anticonvulsant; ophthalmological;
KW	antirheumatic; antiarthritic; antididiabetic; antipsoriatic;
KW	antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;
KW	acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;
KW	neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease;
KW	gene; ds.
XX	
OS	Homo sapiens.
XX	
PN	WO200200928-A2.
XX	
PD	03-JAN-2002.
XX	
PP	02-JUL-2001; 2001WO-EP07537.
XX	
PR	30-JUN-2000; 2000DE-1032529.
XX	
PR	01-SEP-2000; 2000DE-1043826.
XX	
PA	(EPIG-) EPIGENOMICS AG.
XX	
PI	Olek A, Piepenbrock C, Berlin K;
XX	
DR	WPI; 2002-130909/17.
XX	
PT	Nucleic acid comprising fragment of chemically modified gene, useful
PT	for diagnosis and treatment of diseases associated with abnormal
PT	cytosine methylation -
XX	
PS	Claim 1; SEQ ID NO 308; 32pp + Sequence Listing; German.
XX	
CC	The present invention provides a number of human immune system associated
CC	genes which are modified by the methylation of cytosines. The sequences
CC	can be used in the diagnosis and treatment of immune system disorders,
CC	including eye diseases such as retinopathy, neovascular glaucoma and
CC	macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid
CC	leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,
CC	rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel
CC	diseases. The present sequence is a gene of the invention.
XX	
SQ	Sequence 10528 BP; 2873 A; 86 C; 2164 G; 5405 T; 0 other;
	Query Match 29.9%; Score 668.2; DB 24; Length 10528;
	Best Local Similarity 80.2%; Pred. No. 2.6e-175;
	Matches 784; Conservative 0; Mismatches 193; Indels 0; Gaps 0
QY	3 ATTGAACGAGCGATTTCGCCGTATCATCTCACCAACATGCTGTCCACATCTCGTTCTCG 62
Dd	2187 ATTAACAACAAACGGATTTCGCCATATCATCTCACAACATACTATCCACATCTCGTTCTCG 2128
QY	63 GTTATCAGAAATACCAACGAGCGGTGAAGAAGTCACCACCTTTTTTGATGATGATTA 122
Dd	2127 ATTTATCAAAAATACCAACGAAAAACGATAAAAAAATCACCACTTTTTTAATATAATTA 2068
QY	123 CGTGCTCCCTGCATATAATTTGAGCTGAAGCAAATTTGGGCCCAACTCCTGCTCCGCT 182
Dd	2067 CGATACTCCCTATCATATAATTTAACGTAACAAACATATAAACCCAACCTCCTACCTCCGCT 2008
QY	183 CTACTCGTGGTGTTTCATCTTTGGTTTTGGGGCAACATGCTGGTGGTCTCATCTTAAT 242
Dd	2007 CTACTCGTAAATTCATCTTTAAATTTTATAACAACATACTAACTGCTCATCTTAAT 1948
QY	243 AAAC TGCAAAAAGCTGAAGTGCTTGACTGACATTTACCTGCTCAACCTGGCCATCTCTGA 302
Dd	1947 AAAC TACAAAAAAGCTAAAAATACTTAAC TAACATTTTACCTACTCAACCTAACCATCTCTAA 1888
QY	303 TCTGCTTTTTTCTTATTACTCTCCCATTGTGGGTCTACTCTGCTGCAAAATGAGTGGGTCTT 362
Dd	1887 TCTACTTTTTTCTTATTACTCTCCCATTTAAACTCACTCTACTTACAAATAAATAAATCTT 1828

PR	30-JUN-2000; 2000DE-1032529.
FR	01-SEP-2000; 2000DE-1043826.
XX	(EPIC-) EPIGENOMICS AG.
PA	Olek A, Piepenbrock C, Berlin K;
XX	WPI; 2002-130909/17.
DR	Nucleic acid comprising fragment of chemically modified gene, useful
XX	for diagnosis and treatment of diseases associated with abnormal
PT	cytosine methylation -
PT	
PS	Claim 1; SEQ ID NO 307; 32bp + Sequence Listing; German.
XX	
CC	The present invention provides a number of human immune system associated
CC	genes which are modified by the methylation of cytosines. The sequences
CC	can be used in the diagnosis and treatment of immune system disorders,
CC	including eye diseases such as retinopathy, neovascular glaucoma and
CC	macular degeneration, arteriosclerosis, anemia, cancer, acute myeloid
CC	leukemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,
CC	rheumatoid arthritis, psoriasis and inflammatory/alicerative bowel
CC	diseases. The present sequence is a gene of the invention.
XX	
SQ	Sequence 10528 BP; 3072 A; 86 C; 2419 G; 4951 T; 0 other;
Query Match 28.7%; Score 639.8; DB 24; Length 10528;	
Best Local Similarity 78.0%; Pred. No. 2.2e-167;	
Matches 770; Conservative 0; Mismatches 217; Indels 0; Gaps 0;	
QY	1 GGATTGACAAAGGAGCGCATTTCCCAGTACATCCACAACATGCTGCACATCCTGGTCT 60
DB	
DB	8340 GGATTGATAAGGAGCTAATTTTTTAGTATATTTATATATATGTGTTATATATTCGGTTT 8399
QY	61 CGGTTTATCGAANATACCACGAGAGCGGTGAAGAAGTCACCACTTTTTTGATTATGAT 120
DB	
DB	8400 CGGTTTATGAAAATATTAAACGAGAGCGGTGAAGAAGTTATTATTTTTTTGATTATGAT 8459
QY	121 TAGCGTGCTCCCTGTCATAAATTTGACGTGAAGCAAAATGGGCCCAACTCCTGCCCTCG 180
DB	
DB	8460 TAGCGTGTTTTGTTATAAATTTGACGTGAAGTAATATGGGGTTTAAATTTTTGTTTCG 8519
QY	181 CTCFACGCGCGGTGTTTCATCTTTGGTTTGTGGGCAACATGCTGCTGCTCATCTTA 240
DB	
DB	8520 TTTTATTCGTTGGTGTTTATTTTGGTTTGTGGGTAAATGTTGGTCGTTTTTATTTTA 8579
QY	241 ATAACTGC AAAAGCTCAAGTGTCTTCACTGACATTTACCTGCTCAACCTGGCCATCTCT 300
DB	
DB	8580 ATAATTGTAAAAAGTTGAAGTGTTTGATTGATATTTATTTGTTTAAATTTGGTTATTTT 8639
QY	301 GATCTGCTTTTCTTATTACTCTCCCATTTGGGCTCACCTCTGCTGCAAAATGAGTGGGTC 360
DB	
DB	8640 GATTTGTTTTTTTTTATTATTATTTTATTGCGGTATTATTTGTTGTAATGAGTGGGTT 8699
QY	361 TTGGGAATGCAATGTCGCAATTTATCCAGAGGCTGTATCACATPCGGTTATTTGGCGGA 420
DB	
DB	8700 TTGGGAATGTAATGTGTAATATTATTATAGGTTGTGTTATATATCCGTTATTTTGGCGGA 8759
QY	421 ATCTCTTCATCATCCCTCCGAGCAATCGATAGATACCTGGCTATTGPGCATGCTGTTT 480
DB	
DB	8760 ATTUTTTTTTATTATTTTTTTTGATAATCGATAGATATTGCTTATGTTATGTTGTTT 8819
QY	481 GCYTTRAAAGCCAGGACGGTCACCTTTGGGCTGGTGCAAGTGTGTATCACCTGGTTGGTG 540
DB	
DB	8820 GTTTTAAAAGTTAGSACGGTTATTTTGGGCTGGTGATAAGTGTGATTATTTGTTGGTGTG 8879
QY	541 GCTGTGTTTGCTCTGCTCCAGGAATCATCTTTTACTAAATGCCAGAAAGAACGATCTGTT 600
DB	
DB	8880 GTTGTGTTGTTTTGTTTGTAGGAATATTTTATTAATGTTAGAAAGAAATTTTGTT 8939
QY	601 TATGCTGTGGCCCTTATTTTCCAGGATGGAATTAATTTCCACAAATAATAGAGAAC 660
DB	
DB	8940 TATGTTGTGGTTTTATTTTTTACGAGATGAATTAATTTTTTATATAATATAGAGAAAT 8999
DB	

RESULT 15
ABL32334
ID ABL32334 standard; DNA; 10528 BP.

AC ABL32334:

DT 26-MAR-2002 (first entry)

DE	Human	immune system	associated	gene	SEQ ID NO:	307.
yy						

Human: immune system disease; cytosine methylation; antiasthmatic; antiarteriosclerotic; antiamaic; cystostatic; neurotropic; neuroprotective; anti-HIV; antineoplastic; ophthalmologic; antirheumatic; antiarthritis; antidiabetic; anti-psoriatic; antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia; acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy; neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene: ds.

OS Homo sapiens.

PN WO200200928-A2.

PD 03-JAN-2002.

02-JUL-2001; 2001WO-EP07537.

